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THEORETICAL SIMULATION OF SOLAR SPECTRA
IN THE MIDDLE ULTRAVIOLET AND VISIBLE
FOR ATMOSPHERIC TRACE CONSTITUENT MEASUREMENTS

NASA Langley Research Center
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In previous work under this project^(1,2) the stratospheric NO₂ mixing ratio profile was derived from the data obtained during the 9 February 1977 balloon flight. This work has been extended to the 17 February 1977 balloon flight, from which another NO₂ profile was derived. Typical high sun and low sun scans and their ratios already corrected for Rayleigh scattering are shown in Figure 1, along with synthetic spectra of O₃ and NO₂. The spectral absorption coefficients used for NO₂ and O₃ have been tabulated previously⁽¹⁾. Figure 2 shows the NO₂ amounts at three peak absorption wavelengths as determined from several pairs of high sun and low sun spectra. The NO₂ amounts obtained from the two flights are compared in Figure 3 with an earlier NO₂ profile obtained from infrared solar spectra obtained during sunset.⁽³⁾ Only minor changes can be observed among these three winter profiles.

The data obtained during the 9 February 1977 flight (float altitude ~40 km) have also been studied for possible ClO identification. Figure 4 shows high sun and low sun spectra with a simulation of O₃ and ClO absorptions in the 2950-3100A region which are based on the known O₃ absorption coefficients and on recently calculated ClO cross-sections.⁽⁴⁾ The O₃ amount is based on a ray-tracing calculation with a standard O₃ profile;⁽¹⁾ the ClO amount was selected to show a ~10% absorption near 3035A. The ClO absorption increases toward the short wavelength end of Figure 4 and peaks near 2800A. Unfortunately, because the increasing O₃ absorption toward the short wavelength end dominates the spectrum in this region, ClO cannot be identified in these spectra. A similar study is being conducted for the 17 February 1977 flight.

The data compiled during the present study also allow a detailed examination of the attenuation of the solar radiance by the combined effect of the atmospheric trace gases and the SAGE filters. The case of the 0.447μm channel is presented in Figure 5. The amounts of NO₂, O₃ and the airmass were chosen to closely simulate a low sun spectral scan from the 9 February 1977 balloon flight.^(1,2) The solar radiance outside the earth's atmosphere was determined from the AFGL Sacramento Peak Observatory (SPO)⁽⁵⁾ spectral data tape degraded to 2.0A resolution and normalized to Thekaekara's absolute intensities (as described in Ref. 5). The spectral absorption coefficients used for NO₂ and O₃ have been tabulated previously.⁽¹⁾

From these calculations, effective extinction coefficients (weighted by filter response function) can be established which will allow Beer's law (also weighted by the filter response function) to be applied to the analysis of data collected with the 0.447 μ m channel. The coefficients are 3.98×10^{-3} (atm⁻¹ cm⁻¹) and 12.3 (atm⁻¹ cm⁻¹) for O₃ and NO₂ respectively, and 0.229 (airmass⁻¹) for Rayleigh scattering. These calculations will be redone once the final data on this channel become available. The SPO tape provides solar spectral data between 0.38 and 0.7 μ m, so that these computations can be readily performed for the 0.6 μ m channel. The SPO data, though, does not cover the full range of the 0.385 μ m channel or any part of the 1.0 μ m channel. High sun scans from the 9 February 1977 balloon flight can be used to provide the solar spectral data for the 0.385 μ m channel. If calculations of this type are required at 1.0 μ m, an additional source of solar spectral data must be found.

Absorption coefficients have been compiled from the literature^(6,7) for NO₃ in the 4000-7000A region, as shown in Fig. 6. An absorption spectrum has been calculated for 0.001 cm-atm NO₃ as shown in Fig. 7. The NO₃ amount was chosen to produce ~10% absorption near the 6000A peaks. The wavelength region of the stronger peaks were not covered during the February 1977 flights. Comparisons with the flight data showed no observable features due to NO₃. Since the wavelength of the strongest peak (near 6700A) was not covered, and since previous observation showed atmospheric NO₃ during nighttime⁽⁸⁾, it is not surprising that NO₃ was not observed on the current data.

Absorption coefficients have also been compiled for HNO₂⁽⁶⁾, in the 3000-4000A region, as shown in Fig. 8. An absorption spectrum of 0.035 cm-atm HNO₂ is shown in Fig. 9, chosen to produce ~10% absorption near 3700A. No observable features due to HNO₂ could be identified on the Feb. 1977 flights data.

Acknowledgment is made to James Arnold, NASA Ames Research Center for the magnetic tape with the calculated ClO cross-sections. Part of the analysis and the computer work was done by Frederick Fernald and Darwin Rolens. The figures were prepared by Carolyn Bauer. Acknowledgment is made to the National Center for Atmospheric Research, which is sponsored by the National Science Foundation, for computer time used in this research.

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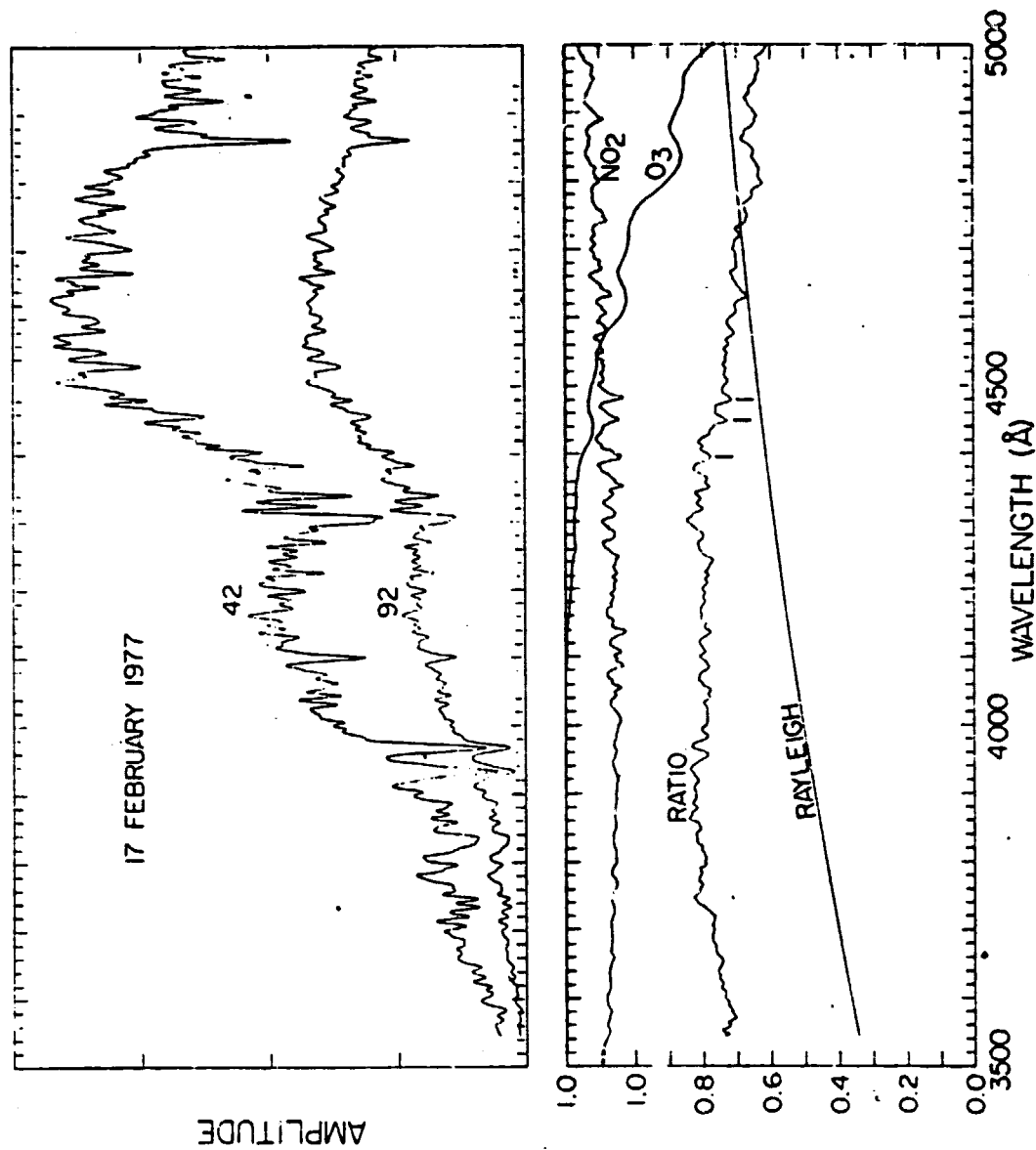


Figure 1. Top Frame: Scans 92 and 42, degraded to $\sim 5\text{\AA}$ resolution. Bottom Frame: ratio of degraded scans 92 to 42, corrected for Rayleigh scattering and synthetic spectra of NO_2 (0.01 atm cm) and O_3 (20 atm cm). The vertical axis for NO_2 and O_3 is transmittance (0 to 1 scale) and is shifted upwards two divisions from the 0 to 1 scale for Rayleigh corrected ratio.

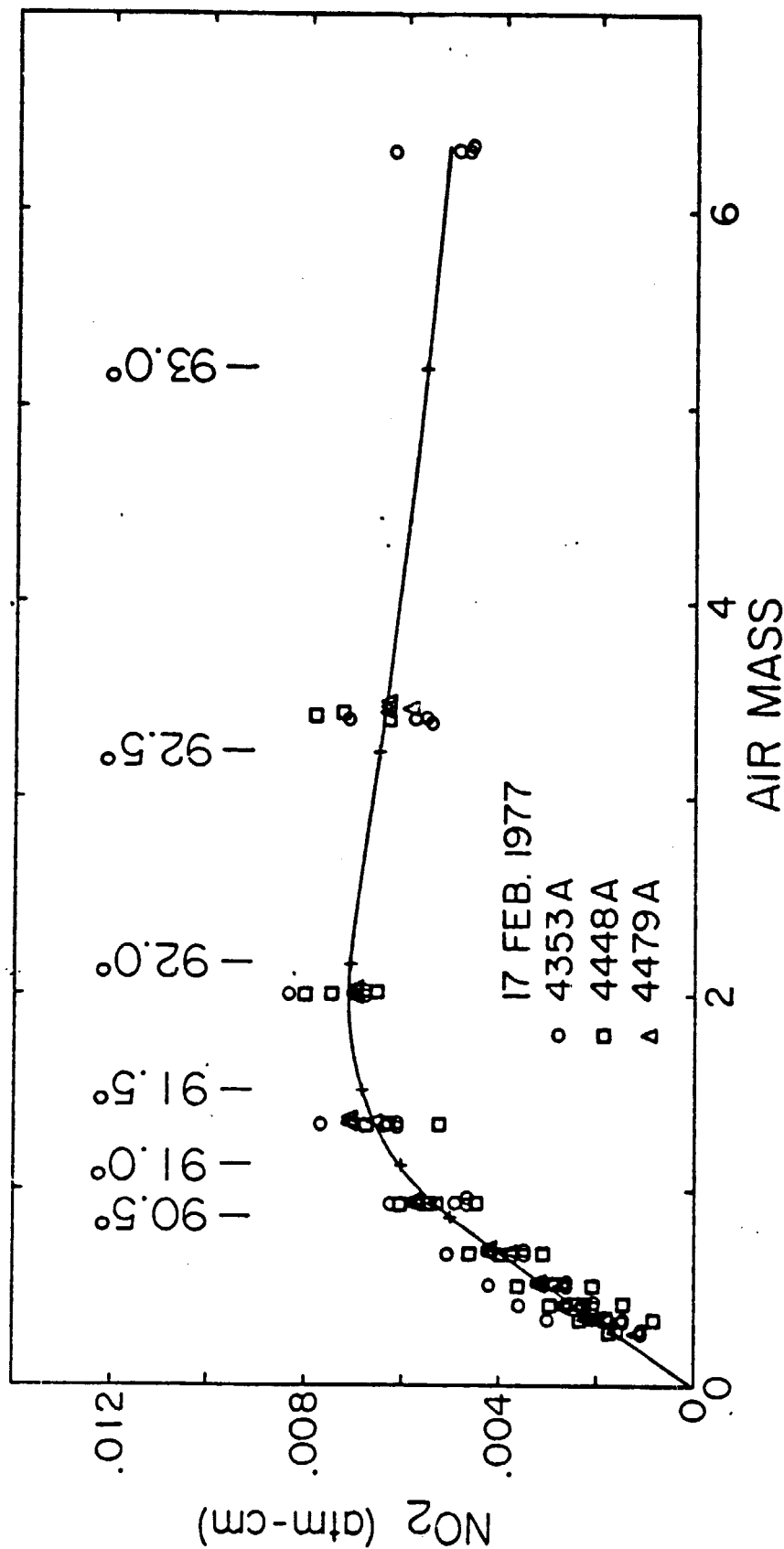


Figure 2. NO₂ amounts and apparent zenith angle as derived from several sunset paths during the 17 February 1977 flight from float altitude of ~30 km. For each of the NO₂ absorption peaks at 4393, 4448 and 4479A, several pairs of high and low sun scans were used. The airmass is calculated at the time corresponding to NO₂ absorption within each scan.

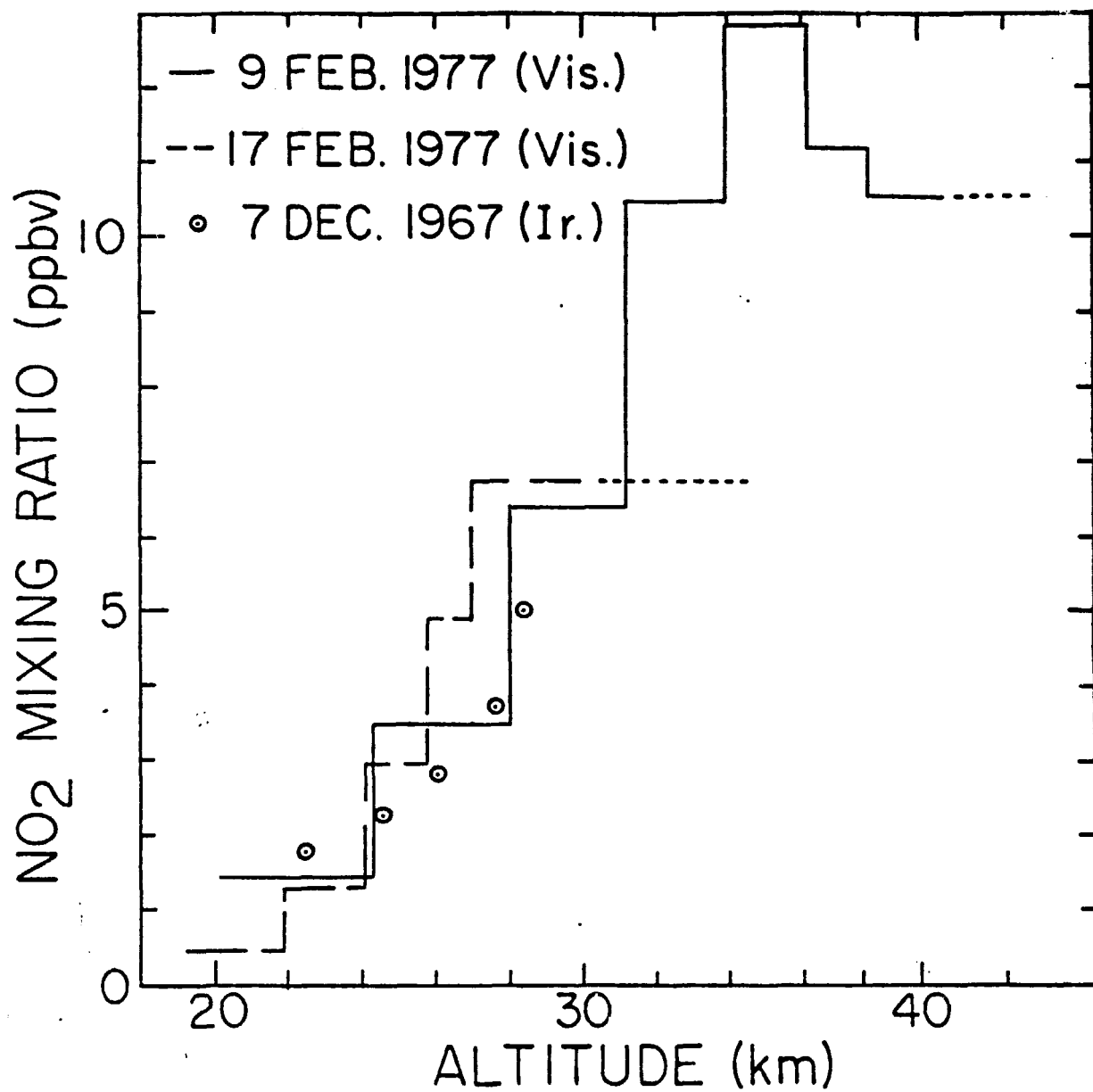


Figure 3. Mixing ratios of NO₂ as derived from three balloon flights from Holloman Air Force Base, New Mexico.

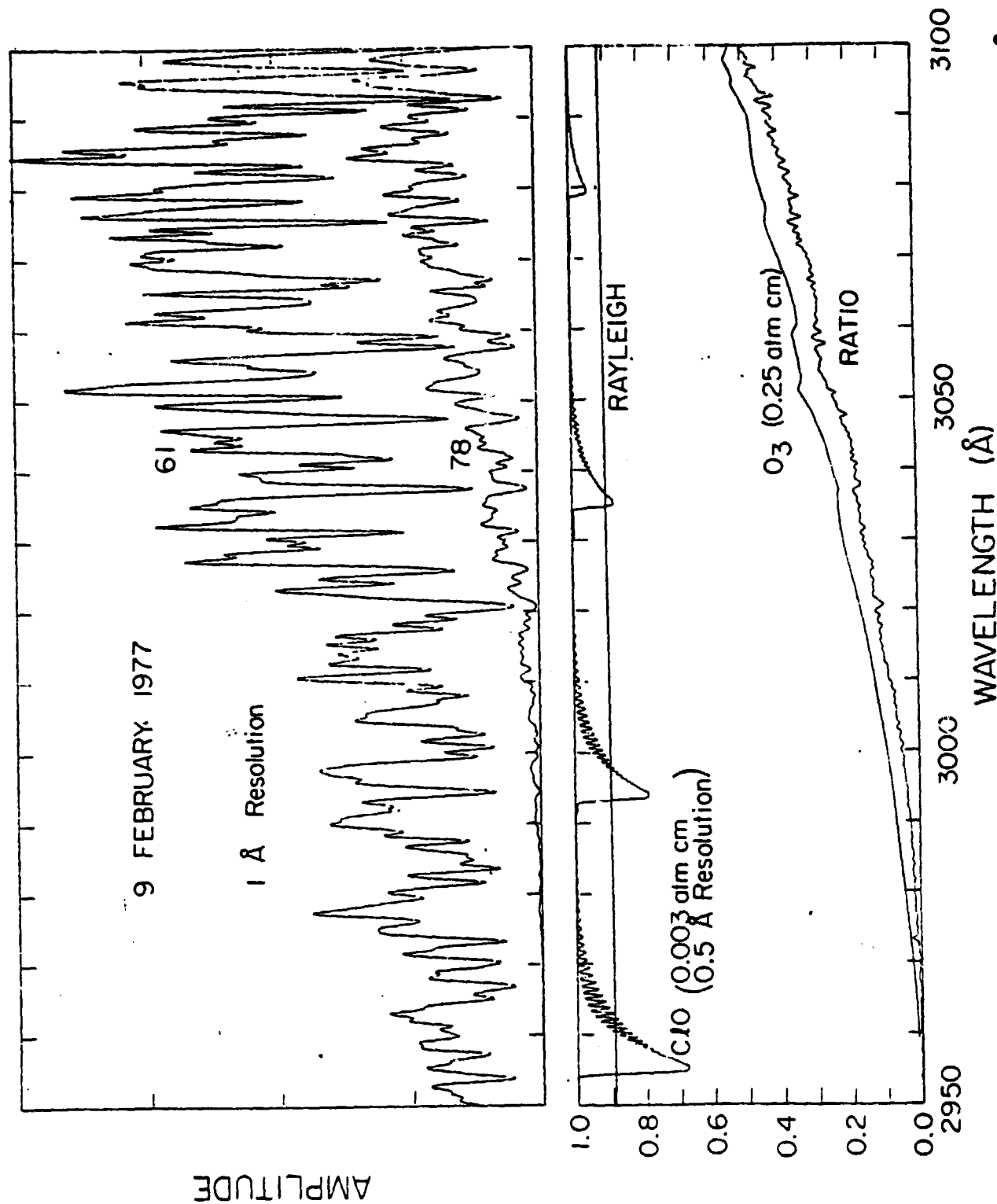


Figure 4. Top Frame: amplitude vs wavelength for scans 61 (solar zenith angle of 78.89°) and 78 (solar zenith angle of 89.02°) from the 9 February 1977 flight degraded to ~1 Å resolution. Bottom Frame: ratio of the degraded scans 78 to 61, corrected for Rayleigh scattering, and synthetic spectrum of 0.25 atm cm O₃ at ~1 Å resolution and 0.003 atm cm O₃ at 0.5 Å resolution. The

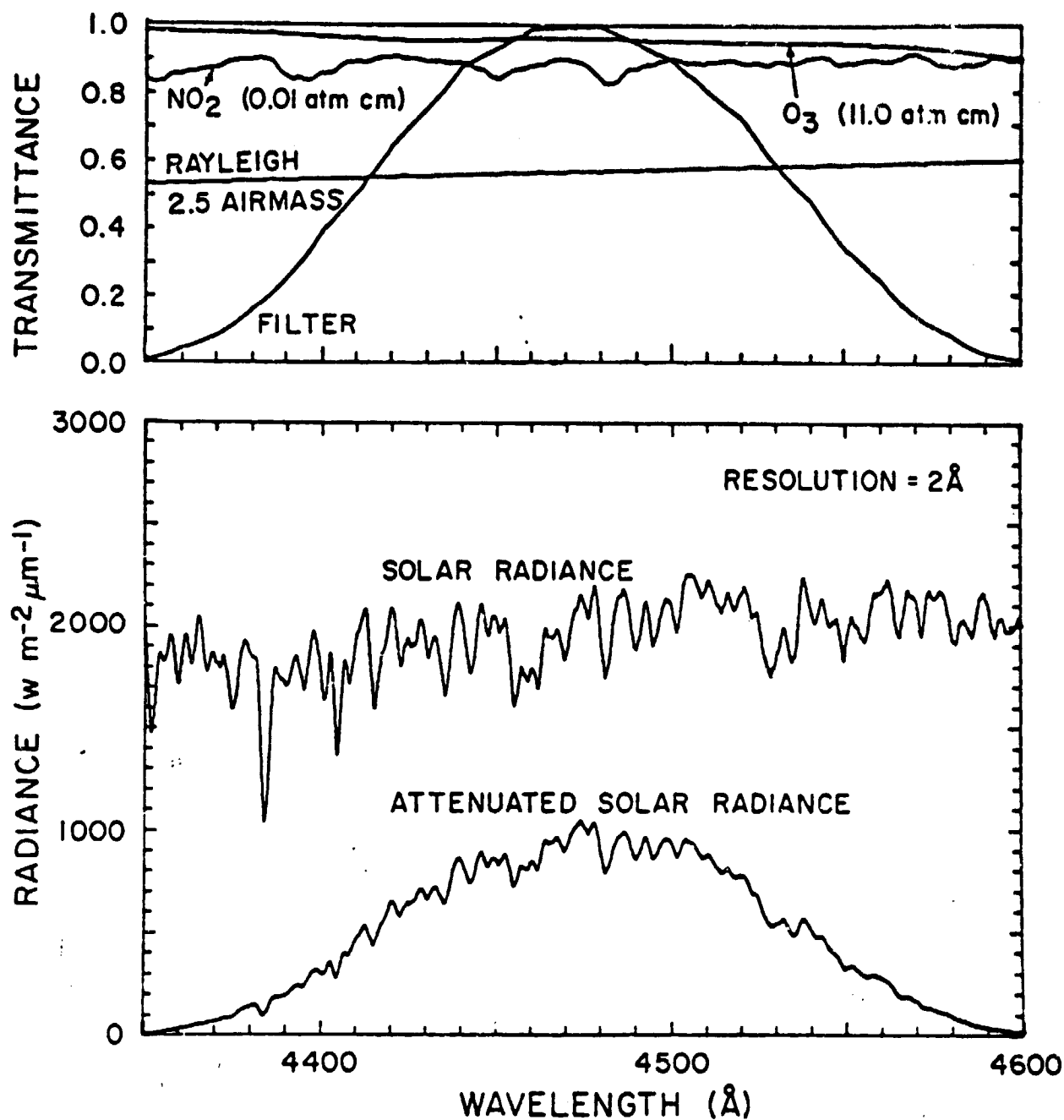


Figure 5. Simulation of the attenuation of the solar spectrum in the 0.447 μ m channel of the SAGE experiment. Top Frame: the normalized filter detector response, the Rayleigh scattering atmospheric transmittance, the NO₂ and O₃ transmittance over the filter bandpass. Bottom Frame: the solar radiance outside the earth's atmosphere as modified by all the transmission losses shown in the top frame.

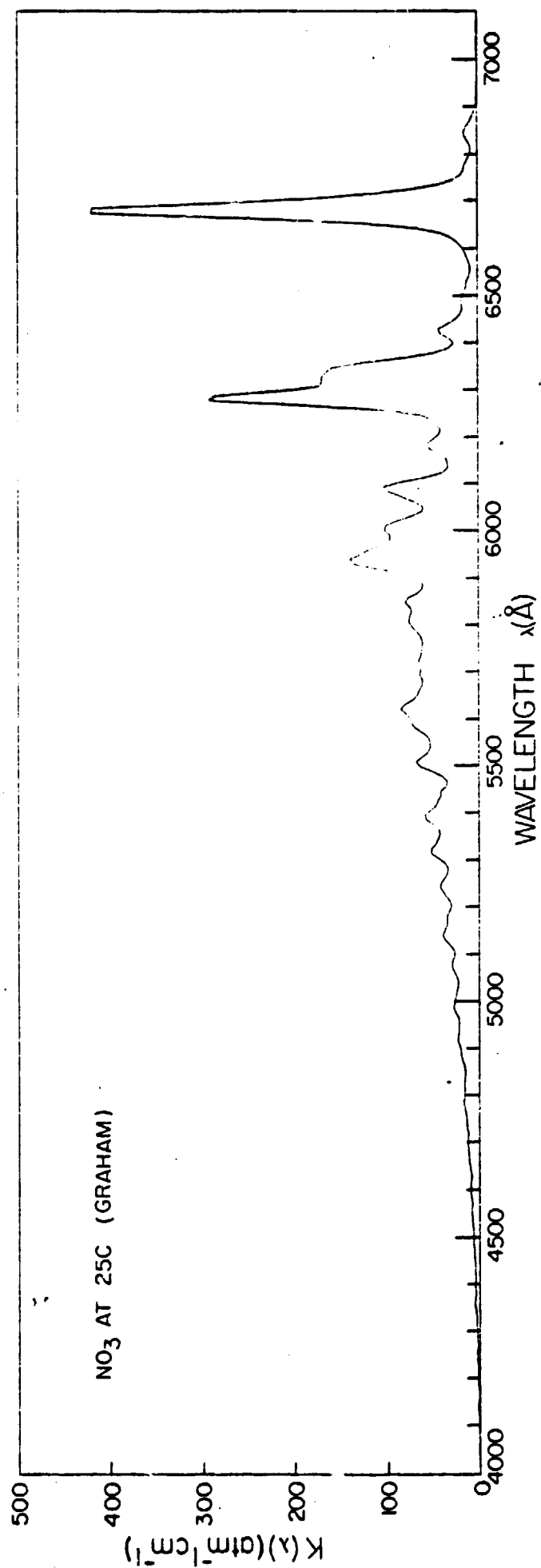


Figure 6. Absorption coefficients of NO_3

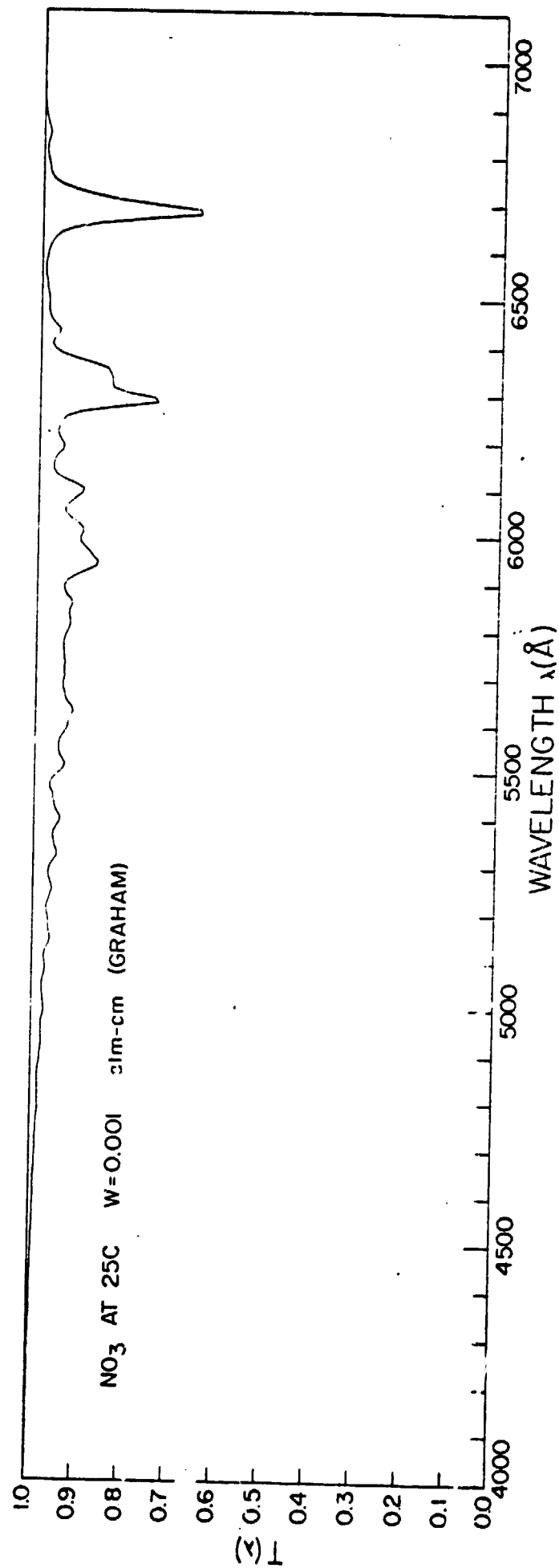


Figure 7. Transmittance of NO₃

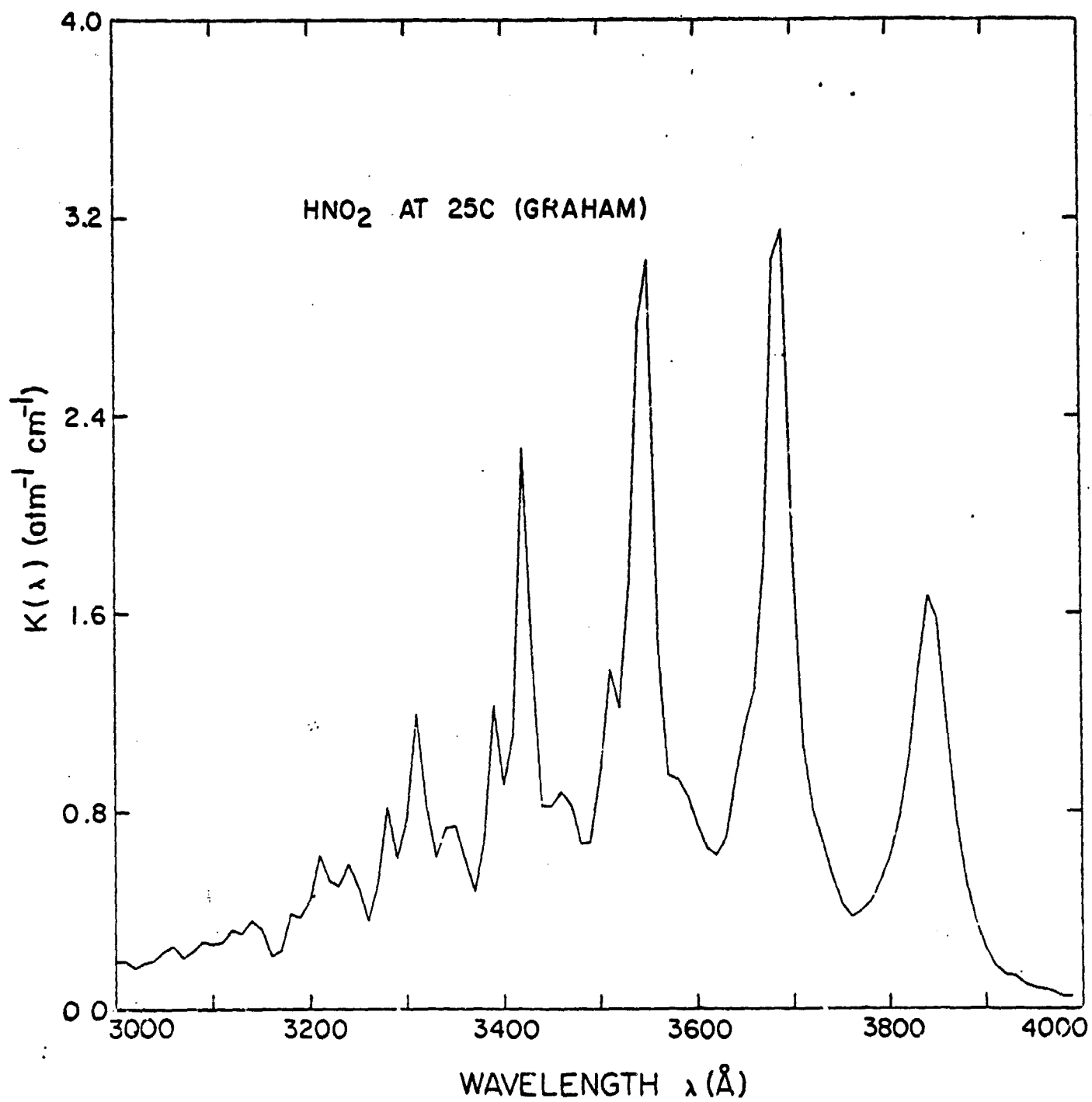


Figure 8. Absorption coefficients of HNO_2

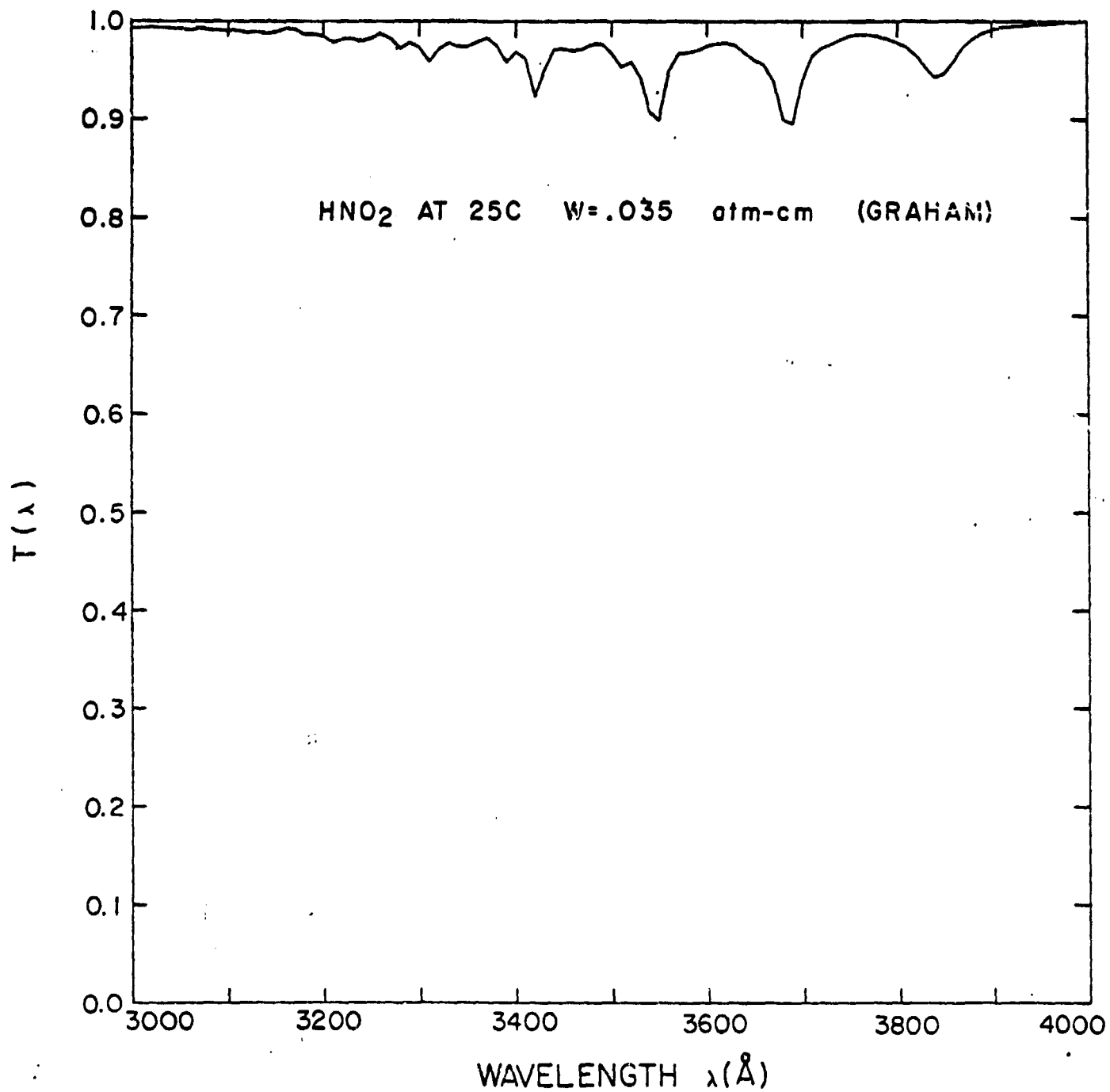


Figure 9. Transmittance of HNO₂